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result set*DB=USPT,PGPB; PLUR=YES; OP=ADJ*

<u>L6</u>	l4 same l3	0	<u>L6</u>
<u>L5</u>	L4 with l3	0	<u>L5</u>
<u>L4</u>	aminopeptidase or IRAP	2499	<u>L4</u>
<u>L3</u>	ptpz or tyrosine phosphatase zeta	31	<u>L3</u>
<u>L2</u>	s ptpz or tyrosine phosphatase zeta	19	<u>L2</u>

DB=USPT,PGPB,JPAB,EPAB,DWPI; PLUR=YES; OP=ADJ

<u>L1</u>	human e2f1	10	<u>L1</u>
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= s ptpz and irap
L1 0 PTPZ AND IRAP

= s aminopeptidase and tyrosine phosphatase zeta
L2 1 AMINOPEPTIDASE AND TYROSINE PHOSPHATASE ZETA

=> dup rem l2
PROCESSING COMPLETED FOR L2
L3 1 DUF REM L2 (0 DUPLICATES REMOVED)

=> d bib ab

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2003 ACS
AN 2000:772846 CAPLUS
DI 133:331185
TI Protein-protein interactions and their use in drug screening and disease
diagnosis
IN Heichman, Karen; Bartel, Paul L.
PA Myriad Genetics, Inc., USA
SO PCT Int. Appl., 87 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000065340	A1	20001102	WO 2000-US10651	20000421
W:	AE, AL, AM, AT, AU, AZ, BA, BE, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KS, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NC, NE, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, EG, KE, MD, RU, TJ, TM				
RW:	GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,				

CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

EP 1181549 A1 20020227 EP 2000-926188 20000421

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, SI, LT, LV, FI, RO

JP 2002542774 T2 20021217 JP 2000-614029 20000421

PRAI US 1999-130389P P 19990422

US 1999-140693P P 19990624

US 1999-156947P P 19990930

US 1999-163073P P 19991102

US 1999-168376P P 19991202

US 1999-168378P P 19991202

WO 2000-US10651 W 20000421

AB The present invention relates to the discovery of novel protein-protein interactions that are involved in mammalian physiol. pathways, including physiol. disorders or diseases. Examples of physiol. disorders and diseases include non-insulin dependent diabetes mellitus, neurodegenerative disorders, such as Alzheimer's disease, and the like. Thus, the present invention is directed to complexes of these proteins and/or their fragments, antibodies to the complexes, diagnosis of physiol. generative disorders (including diagnosis of a predisposition to and diagnosis of the existence of the disorder), drug screening for agents which modulate the interaction of proteins described herein, and identification of addnl. proteins in the pathway common to the proteins described herein.

RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

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